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## The association of circulating inflammatory markers with recurrent vascular events after stroke: a prospective cohort study

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### Abstract

**Background**—Inflammatory markers may be associated with recurrent vascular events after stroke. We aimed to: (1) determine the association between interleukin-6, C-reactive protein, fibrinogen and white cell count and recurrent vascular events after stroke, and (2) compare the association between circulating inflammatory markers and the risk of death from vascular versus non-vascular causes.

**Methods and Results**—We prospectively recruited patients with acute stroke (n=817) and followed them for up to 4 years for the occurrence of: fatal or non-fatal recurrent stroke, myocardial infarction or fatal vascular events, and death from any cause (n=159). The delay to assessment was a median of 10 days. The adjusted incidence of the outcome cluster ‘recurrent stroke, myocardial infarction or vascular death’ after stroke was significantly higher with higher levels of interleukin 6 (75<sup>th</sup> to 25<sup>th</sup> centile: hazard ratio [HR] 1.56, 95 % CI 1.37 to 1.77), C-reactive protein (75<sup>th</sup> to 25<sup>th</sup> centile HR 1.08, 95% CI 1.04 to 1.11) and fibrinogen (75<sup>th</sup> to 25<sup>th</sup> centile HR 1.45, 95% CI 1.24 to 1.72). The associations between inflammatory markers and death were stronger than with recurrent vascular events. The associations of inflammatory markers with vascular and non-vascular deaths were similar.

**Conclusions**—Though inflammatory markers were associated with an increased risk of recurrent vascular events and vascular death after stroke, they were also associated with non-vascular causes of death, suggesting that inflammatory markers do not play a causal role specifically in the generation of recurrent vascular events after stroke. Future studies of the prediction of recurrent vascular events after stroke should concentrate on clinical variables, or different blood markers.

### Keywords

Inflammation; stroke; prognosis

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### DISCLOSURES

None

## INTRODUCTION

In prospective studies of patients with prior stroke or transient ischemic attack (TIA), increased levels of markers of acute inflammation – C-reactive protein (CRP)<sup>1</sup>, interleukin-6 (IL-6)<sup>2</sup>, fibrinogen<sup>1, 3</sup> and white cell count<sup>4</sup> – were associated with increased incidence of recurrent stroke and myocardial infarction (MI) and the risk of death or disability at 6 months after stroke<sup>5</sup>. If high levels of acute-phase markers in the early stages of stroke contribute to recurrent vascular events, we hypothesised the association between inflammatory markers with fatal recurrent vascular events would be stronger than with other, non-vascular, deaths.

In a prospective cohort of patients with recent stroke we aimed to: (1) estimate the association between levels of circulating inflammatory markers and the incidence of ‘recurrent vascular events’ (recurrent stroke, MI and vascular death), and, (2) compare the strength of the association between inflammatory markers and the risk of death from vascular and non-vascular causes.

## METHODS

The Edinburgh Stroke Study was a prospective, hospital-based cohort study of stroke patients followed up for recurrent stroke, MI and death. We have described the methods and process of data collection elsewhere<sup>6</sup>. In brief, we recruited consenting patients with ischemic stroke or intracerebral hemorrhage presenting to the Western General Hospital in Edinburgh between April 2002 and May 2005. We assigned ischemic stroke subtypes (i) according to the site and size of the causative infarct, modified if necessary by imaging findings with the Oxford community stroke project (OCSP) classification and (2) with a modified Trial of ORG10172 in Acute Stroke Treatment (TOAST) classification as described previously.<sup>6</sup> We made a clinical assessment at baseline and contemporaneously drew blood for markers of inflammation (CRP, IL-6, fibrinogen and white cell count and glucose). We obtained follow up data on patients with multiple overlapping methods. We defined as ‘recurrent vascular events’ the outcome cluster ‘recurrent fatal or non-fatal stroke, subsequent fatal or non-fatal MI or other vascular death’. We defined ‘other vascular death’ as deaths due to athero-thrombo-embolic vascular diseases other than stroke or MI. For this study we classified deaths due to the qualifying – though not recurrent - stroke (which were most often due to pneumonia) or gastrointestinal hemorrhage as non-vascular. We did not routinely record the occurrence of infections or other complications between stroke onset and the measurement of vascular outcome or death. The Lothian Research Ethics Committee approved the project.

### Measurement of blood markers

A clinical laboratory measured total white cell count (Beckman Coulter LH750 analyser) and blood glucose (Vitros Chemistry analyser). Blind to clinical details, we measured CRP and fibrinogen in plasma by immunonephelometry (Prospec, Dade Behring Milton Keynes, UK) using the manufacturer's reagents and standards. We assayed IL-6 by ELISA (R & D Systems, Oxford, UK). Intra- and inter-assay coefficients of variation were 4.7 and 8.3%, 2.6 and 5.3%, and 7.5 and 8.9%, respectively.

### Statistical analysis

We used Stata version 10 (Statacorp 2007) for analysis and prepared the paper with reference to the STrengthening the Reporting of OBservational studies in Epidemiology. (STROBE)<sup>7</sup> guidelines. We measured survival to first recurrent vascular event, and separately to each of recurrent stroke, MI and vascular death censoring patients at the end of

follow up or non-vascular death. We compared the baseline characteristics of patients who experienced a recurrent vascular event with those who did not with univariable Cox regression analysis. We examined the relationships between inflammatory markers with correlation coefficients, and used linear regression (after  $\log_e$  transformation of markers) to examine the relationship of markers with delay to blood draw. We used Kaplan Meier survival curves to compare event free survival between groups of patients defined by thirds of inflammatory biomarkers and compared curves with log rank trend tests. We used Cox regression analysis to calculate unadjusted hazard ratios (HR) and 95% confidence intervals (CI) per unit increase in marker levels. We built a multivariate Cox regression model to adjust for confounders, adding variables sequentially that were associated with recurrent vascular events in univariable analysis and had data completeness of over 95%, keeping those variables that significantly improved the fit of the model (likelihood ratio test  $P<0.05$ ). When the model was complete we tested the proportional hazards assumption and its goodness of fit. We looked for first order interactions of inflammatory biomarker levels with other variables in the final model by adding multiplicative terms.

We made further assessment of the marker most strongly associated with recurrent vascular events (IL-6). We assessed the change in discrimination after the addition of IL-6 to a model containing only clinical variables by calculating Harrell's c-statistic for models with and without biomarkers. The c-statistic is analogous to the area under a receiver operator curve for Cox regression models; a value of 0.5 indicates no better discrimination than chance and a value of 1 perfect discrimination.

We replicated the analysis measuring time to death only, censoring at the end of the study. To adjust models examining the risk of death, we used previously validated covariates.<sup>8</sup> We also plotted, by thirds of marker levels, the competing risks of vascular deaths, death due to the initial stroke and death due to other causes using the 'stcompet' command,<sup>9</sup> which calculates the cumulative incidence of each outcome.

## RESULTS

### Baseline characteristics

877 of 1408 patients in the Edinburgh Stroke Study (62%) gave consent and were available for blood to be drawn for markers of inflammation. Of these, 817 (93%) had a definite ischemic stroke, 17 (2%) a probable ischemic stroke and 43 (5%) a hemorrhagic stroke. Of those patients who had blood drawn for blood markers, none was lost to follow up for the outcomes of death, recurrent stroke or myocardial infarction. We first assessed patients clinically at a median of 10 days (IQR 3 to 21 days) after stroke onset and drew blood at a median of 0 days (IQR 0 to 3 days) after assessment. The delay to assessment was longer for outpatients seen in a clinic (median 19 days) than in those admitted to the stroke unit (median 2 days). Patients who did not have blood drawn were similar to those with blood drawn for biomarker data in age, sex distribution and the proportions with hypertension, peripheral or cardiac vascular disease, diabetes or atrial fibrillation. On average, compared to those without blood sampling, patients with blood samples had milder strokes (proportion TACS 6.8 % vs 14.7%  $p=0.001$ ), as patients admitted to hospital and those with more severe symptoms were less likely to be recruited because of practical barriers to obtaining and processing research blood samples and obtaining informed consent or assent.

During the 1866 person years of follow up (mean 2.12 years), 106 patients had a first recurrent stroke (92 ischemic, 5 hemorrhagic and 9 of uncertain type) and 34 had a myocardial infarction. There were 184 deaths: 113 from vascular causes (63 strokes, 35 from cardiac causes, and death from bowel ischemia, vascular dementia and presumed vascular renal failure) and 64 from other causes (33 cancers, 13 chest infections, 6 from

COPD and the rest from pancreatitis, bowel perforation, hip fracture, and extra-pulmonary sepsis).

At the time of the clinical assessment of the index stroke, the median IL-6 was 4.0 (interquartile range [IQR] 2.4 to 7.2) pg/l, median CRP 3.5 (IQR 1.4 to 9.7) mg/l, median fibrinogen 4.5 (IQR 3.8 to 5.4) g/l, median white cell count 8 (IQR 6.6 to 9.7)  $\times 10^9/l$  and median glucose 5.6 (IQR 5 to 6.8) mmol/l. The correlation coefficients were, between IL-6 and other variables were: CRP 0.59, fibrinogen 0.48, glucose 0.06 and white cell count 0.25. The correlation coefficients between  $\log_e$  marker levels and delay from symptom onset to blood draw were, for IL-6  $-0.18$ , CRP  $-0.12$ , fibrinogen  $-0.11$ , white cell count  $-0.13$  and glucose  $-0.06$ . The relationships of average marker levels with the delay from stroke to blood draw were not statistically significant after adjusting for the level of baseline neurological impairment and age.

### Circulating inflammatory markers and recurrent stroke, MI and vascular death

There was a significant increase in recurrent vascular events for patients who: were older; or had a history of AF, heart failure or previous peripheral vascular disease, coronary heart disease or stroke (Table 1). The log hazard of stroke, MI or vascular death rose with each third of IL-6 and CRP, though not by thirds of glucose, fibrinogen or white cell count.

In unadjusted Kaplan-Meier survival analyses, patients survived free of recurrent vascular events for a shorter time in the highest third of IL-6 (log rank trend  $\chi^2=13.2$ ,  $p=0.0003$ ) (figure 1) and CRP (log rank trend  $\chi^2=13.9$ ,  $p=0.0002$ ). This relationship did not reach statistical significance for fibrinogen (log rank trend  $\chi^2=2.8$ ,  $p=0.09$ ), glucose (log rank trend  $\chi^2=1.1$ ,  $p=0.3$ ) or white cell count (log rank trend  $\chi^2=3.1$ ,  $p=0.08$ ). However, a linear model fitted the data well for each marker.

Table 2 shows the association between circulating inflammatory markers and recurrent vascular events. In univariate analyses, all markers except glucose were significantly associated with recurrent vascular events. The relative hazard for an increase of 1 pg/ml of IL-6 was 1.07 (95% CI: 1.04 to 1.10) per pg/ml. The unadjusted associations between IL-6 and recurrent fatal or non-fatal stroke alone (HR 1.04 95% CI 1.00 to 1.08 per pg/ml) were weaker while the HRs for a mg/l increase of CRP, a g/l increase of fibrinogen, a  $1 \times 10^9$  increase in white cell count or a mmol/l increase of glucose were not significantly different from 1. The unadjusted association between IL-6 and fatal or non-fatal MI alone (HR 1.09, 95% CI 1.03 to 1.15) was stronger than for recurrent stroke alone.

We adjusted for the following confounders in the final model: age, prior stroke or TIA or ischemic heart disease, current or prior AF and cardiac failure. Adding markers of stroke severity (i.e. ability to walk or lift arms off bed), blood pressure at assessment, stroke pathological type, diabetes, carotid stenosis, delay to blood taking or smoking did not significantly improve models containing a single inflammatory marker. After adjustment, there was still a significant association between recurrent vascular events and increasing levels of IL-6, CRP and fibrinogen (Table 2). In this cohort, those patients with highest blood levels (75<sup>th</sup> centile) of interleukin 6 had a 1.33 fold increase in the incidence of recurrent vascular events compared with those with the lowest levels (25<sup>th</sup> centile). A similar relative increase in incidence was seen for fibrinogen (HR 1.20), and less for C-reactive protein (HR 1.06).

We added markers sequentially as continuous variables, in order of the strength of their association with recurrent vascular events, to a model containing only clinical variables (age, prior stroke or TIA or heart disease, current or prior AF or cardiac failure). Addition of IL-6 significantly improved the model (likelihood ratio (LR) test  $\chi^2=14.0$ ,  $p<0.001$ ), though

further addition of CRP (LR test  $\chi^2=0.3$ ,  $p=0.56$ ), fibrinogen (LR test  $\chi^2=0.2$ ,  $p=0.68$ ), white cell count (LR test  $\chi^2=0.7$ ,  $p=0.40$ ), or glucose (LR test  $\chi^2=1.3$ ,  $p=0.25$ ) did not make further improvement, probably as the markers were correlated. After adjustment for all markers, only the association between IL-6 and recurrent vascular events remained statistically significant. The final model including IL-6 fulfilled the proportional hazards assumption and fitted the data well.

A model with only clinical variables (age, prior TIA, MI or stroke and AF) had a Harrell's c statistic of 0.62; when we added interleukin-6 to this model, the Harrell's c statistic increased by a small amount, to 0.64.

### Modification of associations by baseline variables

Multiplicative interaction terms between delay to blood taking after stroke, large vessel stroke versus all others (with a modified TOAST algorithm), age, ability to walk and level of IL-6 did not make important changes to the association between IL-6 and recurrent vascular events (and none significantly improved the fit of the final Cox proportional hazards model). This means that, for example for the variable 'delay to blood taking', we were unable to demonstrate a difference in the association between IL-6 and recurrent vascular events in those where blood was taken early (<5 days) or later (>5 days) after stroke. There were no significant two way interactions between other blood markers and IL-6.

### Circulating inflammatory markers and death

All markers were significantly associated with an increased risk of death (Table 3). After adjustment for factors which are known reliably to influence survival after the index stroke (age, being able to walk or talk, independence of daily activities prior to stroke, being able to lift arms from the bed), these associations remained statistically significant though attenuated. IL-6, CRP, fibrinogen and glucose were more strongly associated with death than with recurrent vascular events, though white cell count was less strongly associated. After additional adjustment for all other markers, only the associations of IL-6 and fibrinogen with death remained statistically significant. The association between higher levels of IL-6, CRP and fibrinogen and an increased incidence of death was consistent for each of the separate causes of death (vascular deaths, deaths due to the initial stroke and deaths due to other causes) (figure 2, data for CRP and fibrinogen not shown). Where the cause of death was the qualifying stroke, patients in the top third of the IL-6 distribution had the shortest survival time.

## DISCUSSION

In this cohort consisting of stroke patients on the stroke unit assessed soon after onset and of outpatients with milder strokes seen after a short interval, higher levels of IL-6, CRP and fibrinogen were associated with a higher incidence of recurrent stroke, MI or vascular death, independent of atrial fibrillation, prior vascular events and age. In addition, higher levels of each inflammatory marker were associated with a higher incidence of death from all causes, an association that was stronger than for all vascular events. The association with recurrent stroke alone was weak for IL-6, and did not reach statistical significance in this cohort for the other markers.

We found no consistent evidence of different strengths of association between higher levels of IL-6 with large vessel type strokes versus other stroke subtypes, strokes of different severity or different times from stroke onset to blood draw. Stroke patients had qualitatively similar associations between IL-6, CRP and fibrinogen and deaths from vascular and from



non vascular causes. However, there was a suggestion that early deaths from the index stroke might be more strongly associated with higher levels of inflammation.

### Strengths and limitations

This study had a number of methodological strengths: the cohort included both mild and severe strokes; we used several overlapping methods to ensure we detected all recurrent vascular events; we determined vital status at the end of the follow up period for all of the cohort; data on all suspected outcome events were checked by the study clinicians, either directly or by review of the medical and imaging records. The majority of patients with recurrent strokes underwent repeat brain imaging (93%).

We were unable to draw blood for inflammatory markers from all patients. The most common reasons for not drawing blood were: either the patient did not consent or the practical constraints in inpatients (chiefly the working hours of research laboratories handling the samples). Patients without blood samples tended to have more severe strokes though were otherwise similar (there was no evidence of an interaction between stroke severity and the association between inflammatory markers and either recurrent vascular events or death). These selection biases may have influenced the results.

We drew blood as soon as possible after assessment (median 2 days among patient admitted to hospital); hence levels of IL-6 and CRP were higher than in previous studies (delay to blood draw in these studies was between 12 hours and 30 days), possibly due in some cases to stroke complications such as pneumonia or deep vein thrombosis. As we were unable to adjust for these in our analysis, the observed association may have been due to confounding by these complications. However, in the 60% of patients seen as outpatients, who had milder strokes, and probably fewer infections or other complications, the median time to blood draw was 19 days. Despite this, we were unable to demonstrate effect modification by the time to blood draw after stroke on the association between either IL-6 or CRP and recurrent vascular events, though statistical tests for effect modification are of low power. It is also possible that among patients in whom the initial assessment was delayed, we may have overlooked some early recurrent strokes.

We did not find that large vessel stroke subtypes had a stronger association between marker levels and recurrent events. However, a relatively large number of strokes were unclassified by the TOAST classification, so we cannot exclude the possibility that an association exists.

We used a competing risks survival analysis to examine the association of IL-6, fibrinogen and CRP with of the three main causes of death; vascular, non-vascular and deaths due to the initial stroke. It is possible that there was some misclassification of the cause of death, particularly for deaths occurring soon after stroke when accurate attribution of the cause of death is difficult, even if autopsy is performed.

The epidemiological association between inflammatory markers and recurrent vascular events appears consistent and strong, and similar for IL-6 and CRP, though a somewhat weaker for fibrinogen. However, the degree of extra clinical utility obtained by adding an inflammatory marker to a clinical predictive model is not merely determined by the fact that it is a statistically independent predictor in multivariate models. The small increase in the c-statistic achieved by adding IL-6 to a model based upon clinical variables makes it unlikely that it will add clinically useful prediction to models based on variables that do not require blood draw. We did not calculate a threshold of IL-6 that best predicted recurrent vascular events, as such calculations usually lead to a biased assessment of the strength of predictive ability of markers and are unlikely to be replicated in validation cohorts.<sup>10</sup>

## Interpretation

These data suggest that CRP or IL-6 may not have a major causal role in recurrent vascular events after stroke. It is more likely that the observed association reflects an inflammatory response either to atherosclerosis or to its risk factors, or to an as yet unidentified trigger. In support of this, studies that have examined functional CRP and IL-6 polymorphisms (which produce differences in baseline CRP or IL-6 levels) found different polymorphisms were not associated with an increased risk of stroke<sup>11</sup> or other occlusive vascular events<sup>12</sup>. However, to determine reliably any causal relationship between the physiological inflammatory response and recurrent vascular events in humans would require a randomised trial of an agent which was an effective anti-inflammatory but had no direct effect on other vascular risk factors such as cholesterol or blood pressure.

## Generalisability

Our finding that IL-6 showed the strongest association with recurrent vascular events and with death in this cohort of stroke patients, is consistent with recent reports from population-based prospective studies.<sup>13 14</sup> Our estimates of the association between: (i) CRP, IL-6, fibrinogen and recurrent stroke,<sup>1, 15 16</sup> and (ii) CRP, fibrinogen and death<sup>16, 17</sup> are consistent with previous studies (supplementary table 1).

## Conclusions

We have demonstrated an association between higher levels of IL-6, CRP and fibrinogen and an increased incidence of occlusive vascular events in patients after stroke. The association between IL-6, CRP and fibrinogen and fatal vascular and non-vascular events after stroke seems similar. Future studies of prediction of recurrent vascular events after stroke might better study easily measured clinical variables, or examine the effect of different blood markers.

## Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

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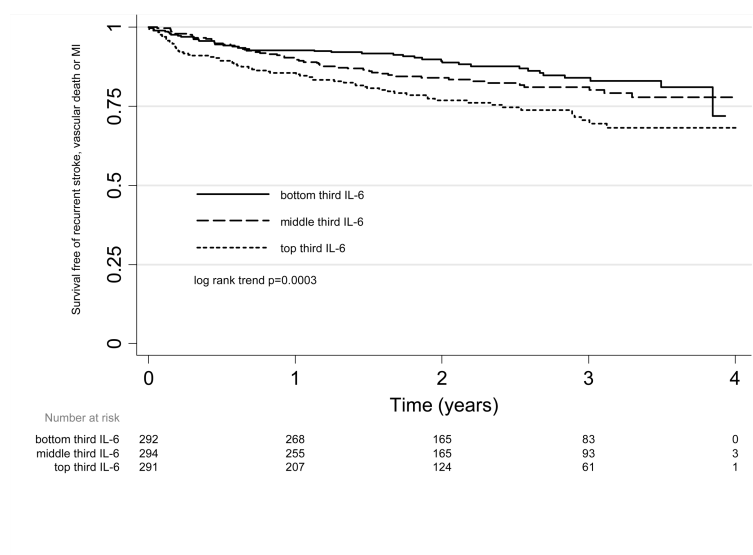
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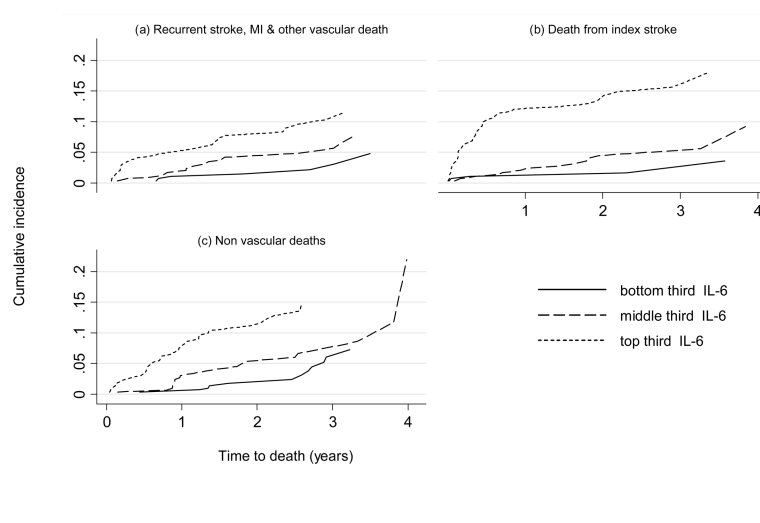
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**Figure 1.** Unadjusted Kaplan Meier survival curve and life table, for survival free from recurrent stroke, myocardial infarction or vascular death by third of interleukin 6.



**Figure 2.** Unadjusted cumulative incidence curves of (a) death from recurrent stroke, MI or other vascular causes, (b) death from the initial stroke with no evidence of recurrent stroke or (c) other non-vascular deaths, by thirds of interleukin 6, estimated from a competing risk analysis.

**Table 1**

Baseline characteristics of stroke patients.

	Total	Recurrent Stroke, MI or vascular death	No vascular event	Univariate HR (95% CI)
<b>Number</b>	877	159	718	
<b>Demographic</b>				
Age, years (mean, SD)	71.4 (12.0)	73.6 (10.7)	70.9 (12.2)	1.02 (1.00 – 1.04) <sup>‡</sup>
Male sex, N (%)	463 (52.8)	80 (50.3)	383 (53.3)	0.9 (0.6 to 1.2)
<b>Laboratory results</b>				
	<b>Median (IQR)</b>	<b>Median (IQR)</b>	<b>Median (IQR)</b>	
Interleukin-6 (pg/ml)	4.0 (2.4-7.2)	4.8 (2.8-9.1)	3.8 (2.3-6.6)	
CRP (mg/L)	3.5 (1.4-9.7)	5.9 (1.9-15.5)	3.3 (1.2-8.8)	
Fibrinogen (g/L)	4.5 (3.8-5.4)	5.7 (3.9-5.7)	4.4 (3.8-5.4)	
White cell count ( $\times 10^9/l$ )	8 (6.6-9.7)	8.4 (6.7-9.7)	7.9 (6.6-9.7)	
Glucose (mmol)	5.6 (5-6.8)	5.7 (4.9-7.2)	5.6 (5-6.7)	
Cholesterol (mmol/l)	5.1 (4.4-6.0)	4.9 (4.3-5)	5.1 (4.4-6)	
<b>Pathological type, index stroke</b>				
	<b>N (%)</b>	<b>N (%)</b>	<b>N (%)</b>	
Definite ischemic	817 (93.2)	149 (93.7)	668 (93.0)	1.1 (0.6 to 2.1) <sup>‡</sup>
Definite hemorrhagic	43 (4.9)	9 (5.7)	34 (4.7)	1.2 (0.6 to 2.4) <sup>‡</sup>
Pathological type unknown	17 (1.9)	1 (0.6)	16 (2.2)	0.3 (0.1 to 1.9) <sup>‡</sup>
<b>Clinical stroke syndrome of index stroke (OCSP<sup>*</sup>)</b>				
TACI	57 (6.8)	10 (6.3)	53 (7.4)	1.2 (0.6 to 2.2) <sup>§</sup>
PACI	376 (45.1)	81 (50.9)	308 (42.9)	1.1 (0.8 to 1.6) <sup>§</sup>
LACI	228 (27.3)	38 (23.9)	200 (27.9)	0.8 (0.5 to 1.1) <sup>§</sup>
POCI	131 (15.7)	22 (13.8)	121 (16.9)	0.8 (0.5 to 1.3) <sup>§</sup>
Uncertain subtype	42 (5.0)	8 (5.0)	36 (5.0)	0.9 (0.5 to 1.9) <sup>§</sup>
<b>Severity of index stroke</b>				
Can't walk, can't lift both arms	91 (10.4)	12 (7.6)	79 (11.0)	0.9 (0.5 to 1.5)
Can't walk, can lift both arms	119 (13.6)	31 (19.5)	88 (12.3)	1.6 (1.1 to 2.4)
Can walk	664 (76.0)	115 (72.9)	549 (76.7)	0.7 (0.5 to 1.0)
<b>Aetiological classification of index stroke (TOAST)</b>				
Cardioembolic	117 (13.3)	28 (17.6)	89 (12.4)	1.2 (0.8 to 2.0) <sup>//</sup>
Large vessel disease	72 (8.2)	12 (7.6)	60 (8.4)	1.0 (0.7 to 1.6) <sup>//</sup>
Mixed aetiology	58 (6.6)	16 (7.6)	42 (5.9)	1.8 (1.2 to 2.7) <sup>//</sup>
Small vessel disease	178 (20.3)	24 (15.1)	154 (21.5)	0.8 (0.6 to 1.2) <sup>//</sup>
Unclassified after complete investigation	355 (40.5)	65 (40.9)	290 (40.4)	0.9 (0.7 to 1.3) <sup>//</sup>
Unclassified after incomplete investigation	40 (15.8)	14 (8.8)	83 (11.6)	0.8 (0.5 to 1.5) <sup>//</sup>
<b>Risk factors</b>				
History of TIA	143 (16.3)	29 (18.2)	114 (15.9)	1.1 (0.8 to 1.7)
History of stroke	166 (18.9)	41 (25.8)	125 (17.4)	1.5 (1.1 to 2.2)

	Total	Recurrent Stroke, MI or vascular death	No vascular event	Univariate HR (95% CI)
History of ischemic heart disease	242 (27.6)	62 (39.0)	180 (25.1)	1.9 (1.4 to 2.6)
History of PVD	69 (7.9)	22 (13.9)	47 (6.6)	2.0 (1.3 to 3.2)
Ipsilat. carotid stenosis >70	97 (12.5)	21 (14.9)	76 (12.0)	1.2 (0.8 to 2.0)
Ever AF	168 (20.3)	42 (26.4)	126 (17.6)	1.7 (1.3 to 2.5)
Prior treated hypertension	467 (53.3)	96 (60.4)	371 (51.7)	1.4 (1.0 to 1.9)
Diabetes	110 (12.5)	26 (16.4)	84 (11.7)	1.4 (0.9 to 2.2)
Ever smoker	602 (69.8)	113 (71.1)	489 (69.5)	1.1 (0.8 to 1.5)
Heart failure	40 (4.58)	14 (8.7)	26 (3.6)	2.8 (1.6 to 4.8)
Any antiplatelet at baseline	369 (46.1)	11 (6.9)	33 (4.6)	1.7 (0.9 to 3.1)
Warfarin at baseline	43 (4.9)	32 (4.5)	11 (6.9)	1.5 (0.8 to 2.7)
Systolic BP (Mean, No.observations)	147.2 (874)	147.3 (159)	147.2 (715)	1.00 (0.99 to 1.01) <sup>†</sup>
Diastolic BP (Mean, No. observations)	80.0 (874)	80.1 (159)	80.0 (715)	1.00 (0.99 to 1.01) <sup>†</sup>

\* OSCP=Oxfordshire Community Stroke Project Classification (ischemic and probable), TACS=total anterior circulation stroke, PACS=partial anterior circulation stroke, LACS=lacunar stroke, POCS=posterior circulation stroke.

<sup>†</sup> per unit increase

<sup>‡</sup> vs other pathological types

<sup>\$</sup> vs all others in OSCP classification

// vs all others in TOAST classification.



**Table 2**

The association between marker level and recurrent stroke, MI or vascular death, assuming a linear association between marker level and log hazards

	Hazard ratio per unit increase* in marker (95% CI)			Hazard ratio comparing 75 <sup>th</sup> to 25 <sup>st</sup> centile <sup>†</sup>
	Unadjusted	Adjusted <sup>‡</sup>	Adjusted for all markers <sup>§</sup>	Adjusted <sup>‡</sup>
<b>Interleukin-6 (pg/ml)</b>	1.07 (1.04 to 1.10)	1.06 (1.03 to 1.09)	1.05 (1.01 to 1.09)	1.33 (1.15 to 1.53)
<b>C-reactive protein (mg/l)</b>	1.01 (1.00 to 1.01)	1.01 (1.00 to 1.01)	1.00 (1.00 to 1.03)	1.06 (1.02 to 1.09)
<b>Fibrinogen (g/l)</b>	1.16 (1.05 to 1.28)	1.12 (1.01 to 1.25)	1.02 (0.97 to 1.17)	1.20 (1.01 to 1.43)
<b>White cell count (<math>\times 10^9/l</math>)</b>	1.06 (1.02 to 1.10)	1.05 (1.00 to 1.11)	1.03 (0.97 to 1.09)	1.17 (0.98 to 1.38)
<b>Glucose (mmol/l)</b>	1.04 (0.99 to 1.09)	1.03 (0.98 to 1.08)	1.02 (0.97 to 1.08)	1.06 (0.97 to 1.15)

\* for IL-6 pg/ml; CRP mg/l; fibrinogen g/l; white cell count  $\times 10^9/l$ ; glucose mmol/l

<sup>†</sup> 25<sup>th</sup> and 75<sup>th</sup> percentile respectively for: IL6 2.39 and 7.22 pg/ml; CRP 1.39 and 9.65 mg/l; fibrinogen 3.81 and 5.41 g/l; white cell count 6.6 and  $9.7 \times 10^9/l$ ; glucose 5.0 and 6.8 mmol/l

<sup>‡</sup> Adjusted for confounders: age, cardiac failure, atrial fibrillation (current or past), or prior stroke, TIA, peripheral vascular disease or MI.

<sup>§</sup> adjusted for all confounder in previous column, and other markers.

**Table 3**

The association between marker level and death from any cause, assuming a linear association between marker level and log hazards

	Hazard ratio per unit increase* in marker (95% CI)			Hazard ratio comparing 75 <sup>th</sup> to 25 <sup>th</sup> centile <sup>†</sup>
	Unadjusted	Adjusted <sup>‡</sup>	Adjusted for all markers <sup>§</sup>	Adjusted <sup>‡</sup>
<b>Interleukin-6 (pg/ml)</b>	1.13 (1.10 to 1.15)	1.10 (1.07 to 1.12)	1.07 (1.04 to 1.12)	1.56 (1.37 to 1.77)
<b>C-reactive protein (mg/l)</b>	1.01 (1.00 to 1.01)	1.01 (1.00 to 1.01)	1.00 (1.00 to 1.81)	1.08 (1.04 to 1.11)
<b>Fibrinogen (g/l)</b>	1.37 (1.26 to 1.49)	1.26 (1.14 to 1.40)	1.14 (1.01 to 1.28)	1.45 (1.24 to 1.72)
<b>White cell count (<math>\times 10^9/l</math>)</b>	1.07 (1.02 to 1.12)	1.05 (1.00 to 1.11)	1.03 (0.97 to 1.09)	1.17 (1.00 to 1.37)
<b>Glucose (mmol/l)</b>	1.95 (1.02 to 1.10)	1.06 (1.02 to 1.11)	1.04 (0.99 to 1.09)	1.12 (1.03 to 1.21)

\* for IL-6 pg/ml; CRP mg/l; fibrinogen g/l; white cell count  $\times 10^9/l$ ; glucose mmol/l

<sup>†</sup> 25<sup>th</sup> and 75<sup>th</sup> percentile respectively for: IL6 2.39 and 7.22 pg/ml; CRP 1.39 and 9.65 mg/l; fibrinogen 3.81 and 5.41 g/l; white cell count 6.6 and  $9.7 \times 10^9/l$ ; glucose 5.0 and 6.8 mmol/l.

<sup>‡</sup> Adjusted for confounders: age, ability to walk, living alone, independent prior to stroke, orientated to place time and person, able to lift arms from bed.

<sup>§</sup> adjusted for all confounder in previous column, and other markers